FDA 510(k) Premarket Notification

K830529 BIOFEEDBACK-TRAINING INSTRUMENT

CLEARED

Mar 24, 1983
34 days to decision

K830529 · Product code: **HCC** · Neurology
Source: https://www.510kdatabase.net/k830529/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Feb 18, 1983
Decision date	Mar 24, 1983
Days to decision	34 days
Third-party review	No

APPLICANT

Company	**Relaxomat, Inc.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1983-1995